Following is a letter from GlaxoSmithKline dated January 23, 2007, about an urgent drug recall:

Pharmacy and wholesaler level recall of specific lots of Paxil CR® (Paroxetine Hydrochloride) Controlled Release Tablets

GlaxoSmithKline is recalling the following products **from the retail and wholesale levels for the referenced lot numbers only:**

Product	Strength	Lot Number	NDC	Package Size
Paxil® CR Tablets	12.5 mg	9-6P06	0029-3206-13	Bottle 30's
Paxil® CR Tablets	12.5 mg	14-6P06	0029-3206-13	Bottle 30's
Paxil® CR Tablets	12.5 mg	15-6P06	0029-3206-13	Bottle 30's
Paxil® CR Tablets	12.5 mg	16-6P06	0029-3206-13	Bottle 30's

The above referenced Paxil CR® tablet lots are being recalled because a very small number of bottles may display an incorrect dosage strength (25mg) on the Medical Guidance leaflet which forms part of the outer bottle label. This strength will be at variance with that correctly printed on the base label applied directly to the bottle.

For shipping assistance or questions or questions about the recall process, please contact Stericycle, Inc. at 1-866-324-3735. A check will be issued for the price in effect at the time of purchase. In addition, a service and handling charge will be included based on HDMA guidelines. If you have any medical questions, please contact the GlaxoSmithKline Customer Response Center at 1-888-825-5249 between 8:00 a.m. and 8:00 p.m. ET, Monday through Friday.

This recall is being conducted with the knowledge of the Food and Drug Administration. We appreciate your immediate attention and cooperation. GlaxoSmithKline remains committed to product quality, integrity and patient satisfaction and we sincerely regret any inconvenience this action may cause.